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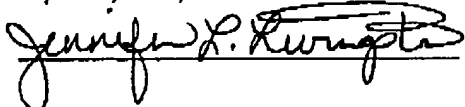
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**FROM: Jennifer L. Livingston**

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**Number of Pages Including this Page: 5**

Listed below are the item(s) being submitted with this Certificate of Transmission:

- 1) Request for Correction of Filing Receipt (1 page)
- 2) Copy of 1<sup>st</sup> page of application listing cross reference information (1 page)
- 3) Filing Receipt with correction noted (2 pages)

Inventor(s): Michael George Natchus et al.

S.N.: 10/730,572

Filed: December 8, 2003

Case: 6229MRR

Comments:

P&amp;G Case 6229MRR

## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In the application of :  
Michael George Natchus, et al. : Confirmation No. 3064  
Serial No. 10/730,572 : Group Art Unit 1614  
Filed December 8, 2003 : Examiner

For Substituted Cyclic Amine Metalloprotease Inhibitors

REQUEST FOR CORRECTION OF FILING RECEIPT

Commissioner for Patents  
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Dear Sir:

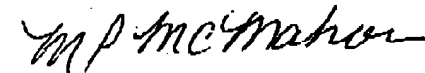
Applicants hereby request a corrected filing receipt. Attached is a copy of the Filing Receipt with the changes noted thereon. Specifically, there is an error in the filing date associated with the first listed cross-reference. The correct filing date for CIP Application No. 10/186,531, as shown on the first page of the filed application (copy attached) is October 1, 2003. The updated filing receipt, mailing date 04/06/2004, lists the date as 07/01/2002. Please make the following correction:

Please omit "07/01/2002" and insert therefore - 10/01/2003 -.

Applicants submit that the error was not the fault of the Applicants. Accordingly, no fee is believed to be due.

Respectfully submitted,

By



Mary Pat McMahon  
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Date: April 21, 2004

Customer No. 27752



## UNITED STATES PATENT AND TRADEMARK OFFICE

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APPL NO.	FILING OR 371 (c) DATE	ART UNIT	FIL FEE REC'D	ATTY. DOCKET NO	DRAWINGS	TOT CLMS	IND CLMS
10/730,572	12/08/2003	1614	900	6229MRR		12	3

CONFIRMATION NO. 3064

27752

THE PROCTER & GAMBLE COMPANY  
INTELLECTUAL PROPERTY DIVISION  
WINTON HILL TECHNICAL CENTER - BOX 161  
6110 CENTER HILL AVENUE  
CINCINNATI, OH 45224

UPDATED FILING RECEIPT



\*OC000000012289842\*

Date Mailed: 04/06/2004

Receipt is acknowledged of this regular Patent Application. It will be considered in its order and you will be notified as to the results of the examination. Be sure to provide the U.S. APPLICATION NUMBER, FILING DATE, NAME OF APPLICANT, and TITLE OF INVENTION when inquiring about this application. Fees transmitted by check or draft are subject to collection. Please verify the accuracy of the data presented on this receipt. If an error is noted on this Filing Receipt, please write to the Office of Initial Patent Examination's Filing Receipt Corrections, facsimile number 703-746-9195. Please provide a copy of this Filing Receipt with the changes noted thereon. If you received a "Notice to File Missing Parts" for this application, please submit any corrections to this Filing Receipt with your reply to the Notice. When the USPTO processes the reply to the Notice, the USPTO will generate another Filing Receipt incorporating the requested corrections (if appropriate).

## Applicant(s)

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CENTRAL DOCKETING	
Atty/GBU Contact: <i>JLL/KLP</i>	
DATE REC'D	APR 8 2004
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## Domestic Priority data as claimed by applicant

This application is a CIP of 10/186,531 07/01/2002 \* 10/01/2003  
which is a CIP of 09/888,675 06/25/2001 PAT 6,569,855  
and is a CIP of 09/888,759 06/25/2001 ABN  
which is a DIV of 08/918,317 08/26/1997 PAT 6,417,219  
which claims benefit of 60/024,842 08/28/1996  
(\*Data provided by applicant is not consistent with PTO records.

## Foreign Applications

If Required, Foreign Filing License Granted: 03/10/2004

Projected Publication Date: 07/15/2004

**Non-Publication Request:** No

**Early Publication Request:** No

**Title**

Substituted cyclic amine metalloprotease inhibitors

**Preliminary Class**

514

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Title 35, United States Code, Section 184  
Title 37, Code of Federal Regulations, 5.11 & 5.15**

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6229MRR

**SUBSTITUTED CYCLIC AMINE METALLOPROTEASE INHIBITORS**

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Biswanath (NMN) De  
Stanislaw (NMN) Pikul  
Neil Gregory Almstead  
Roger Gunnard Bookland  
Yetunde Olabisi Taiwo  
Menyan (NMN) Cheng

**CROSS REFERENCE**

This is a Continuation-in-Part Application of Application No. 10/186,531, filed October 1, 2003, which is a Continuation-in-Part of Application Nos. 09/888,675 and 09/888,759 filed concurrently on June 25, 2001, which are Divisionals of US Application No. 08/918,317, filed August 26, 1997 which claim priority under Title 35, United States Code 119(e) from Provisional Application Serial No. 60/024,842, filed August 28, 1996.

**TECHNICAL FIELD**

This invention is directed to compounds which are useful in treating diseases associated with metalloprotease activity, particularly zinc metalloprotease activity.

**BACKGROUND****Background**

A number of structurally related metalloproteases [MPs] effect the breakdown of structural proteins. These metalloproteases often act on the intercellular matrix, and thus are involved in tissue breakdown and remodeling. Such proteins are referred to as metalloproteases or MPs. There are several different families of MPs, classified by sequence homology. Several families of known MPs, as well as examples thereof, are disclosed in the art.

These MPs include Matrix-Metallo Proteases [MMPs], zinc metalloproteases, many of the membrane bound metalloproteases, TNF converting enzymes, angiotensin-converting enzymes (ACEs), disintegrins, including ADAMs (See Wolfsberg et al, 131 *J. Cell Bio.* 275-78 October, 1995), and the enkephalinases. Examples of MPs include human skin fibroblast collagenase, human skin fibroblast gelatinase, human sputum collagenase, aggrecanase and gelatinase, and human stromelysin. Collagenase, stromelysin, aggrecanase and related enzymes are thought to be important in mediating the symptomatology of a number of diseases.

Many potential therapeutic indications of MP inhibitors have been discussed in the literature. For example, coronary artery disease affects 57.5 million people in the United States, alone, annually claiming the lives of approximately one million individuals. The primary instigator leading to the characteristic sequelae of coronary artery disease is atherosclerosis. Atherosclerosis is a complex interaction between lipids and other elements in the blood, and the